Fig. 1
Study of the release of DNA at pH 5.0

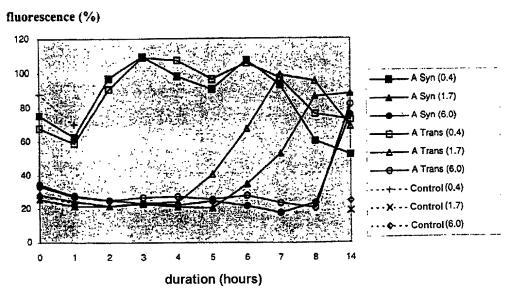
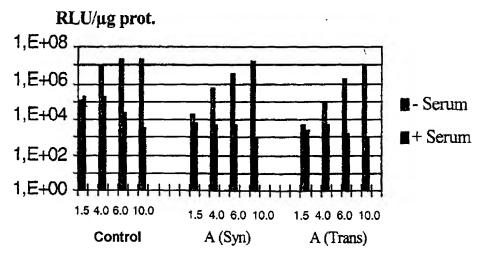


Fig. 2

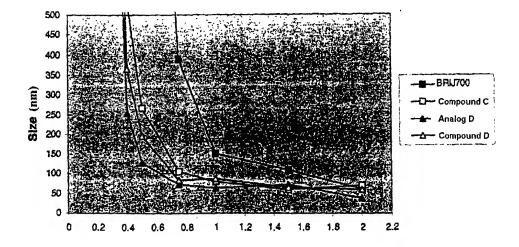
Efficiency of transfection in vitro



DNA/control cationic lipid complexes at different charge ratios

Fig. 3

Stabilization of the nucleolipid complexes by compound C, compound D, BRIJ700 or analog D



Weight/weight ratio with the DNA

Fig. 4

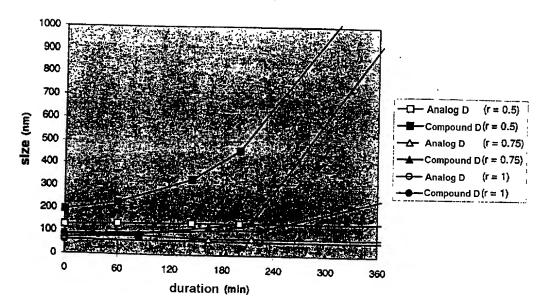
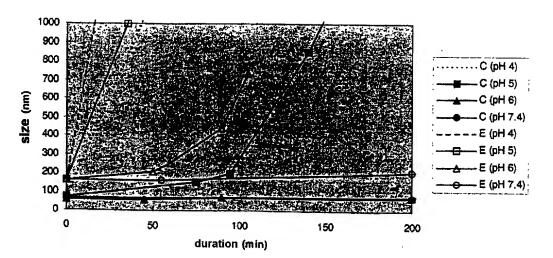
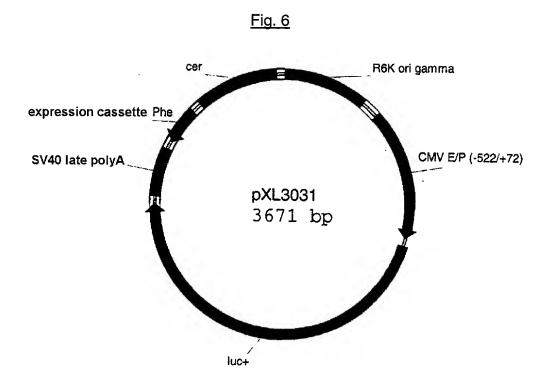


Fig. 5





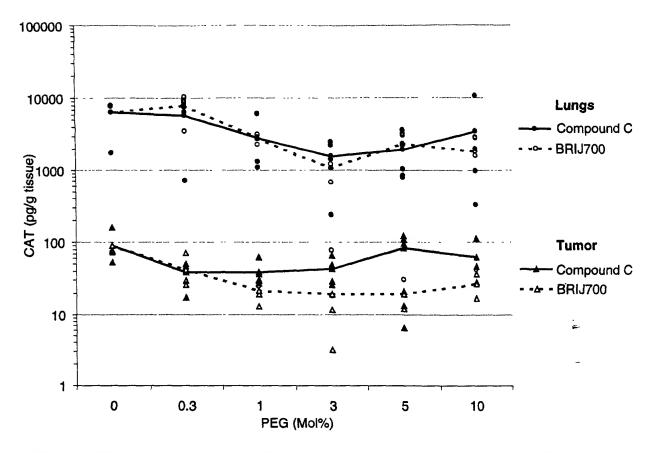


Figure 7: Dose response of pH labile C18-PEG₅₀₀₀ (Compound C) on gene transfer activity in vivo mediated by a cationic lipid/DOPE/DNA (5/5/1) complex. Non-degradable C18-PEG₅₀₀₀ (BRIJ700) was used as a negative control. Data are mean (lines) and individual values of 4 Balb/C mice bearing subcutaneous M109 tumor.

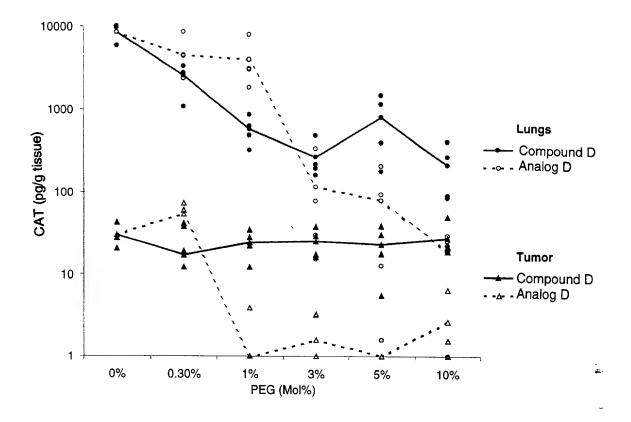


Figure 8: Dose response of pH labile cholesterol-PEG $_{5000}$ (Compound D) on gene transfer activity in vivo mediated by a cationic lipid/DOPE/DNA (5/5/1) complex. Non-degradable cholesterol-PEG $_{5000}$ (Analog D) was used as a negative control. Data are mean (lines) and individual values of 4 Balb/C mice bearing subcutaneous M109 tumor.